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NEWS 9 APR 02 CAS Registry Number Crossover Limits Increased to
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NEWS 13 APR 02 EMBASE Adds Unique Records from MEDLINE, Expanding
Coverage back to 1948
NEWS 14 APR 07 CA/Caplus CLASS Display Streamlined with Removal of
Pre-IPC 8 Data Fields
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NEWS EXPRESS FEBRUARY 15 10 CURRENT WINDOWS VERSION IS V8.4.2,
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Updated Search

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DICTIONARY FILE UPDATES: 3 JUN 2010 HIGHEST RN 1226953-63-4

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=> S 11
SAMPLE SEARCH INITIATED 16:25:40 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 5685 TO ITERATE

35.2% PROCESSED 2000 ITERATIONS 0 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00:00:01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
PROJECTED ITERATIONS: 109179 TO 118221
PROJECTED ANSWERS: 0 TO 0

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L2 0 SEA SSS SAM L1

=> s 11 full
THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 191.05 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y
FULL SEARCH INITIATED 16:25:46 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 114659 TO ITERATE

100.0% PROCESSED 114659 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.07

L3 0 SEA SSS FUL L1

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L4 STRUCTURE UPLOADED

=> s 14
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SAMPLE SCREEN SEARCH COMPLETED - 91 TO ITERATE

100.0% PROCESSED 91 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 1248 TO 2392
PROJECTED ANSWERS: 0 TO 0

L5 0 SEA SSS SAM L4

=> s 14 full
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FULL SCREEN SEARCH COMPLETED - 1807 TO ITERATE

100.0% PROCESSED 1807 ITERATIONS 2 ANSWERS
SEARCH TIME: 00.00.01

L6 2 SEA SSS FUL L4

=> file hcplus
COST IN U.S. DOLLARS SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST 386.02 386.24

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FILE COVERS 1907 - 5 Jun 2010 VOL 152 ISS 24

FILE LAST UPDATED: 4 Jun 2010 (20100604/ED)

REVISED CLASS FIELDS (/NCL) LAST RELOADED: Apr 2010

USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Apr 2010

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2010.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 16
L7 3 L6

=> s 17 and lightner, j?/au
 55 LIGHTNER, J?/AU
L8 0 L7 AND LIGHTNER, J?/AU

=> s 17 and ng, h?/au
 1105 NG, H?/AU
L9 0 L7 AND NG, H?/AU

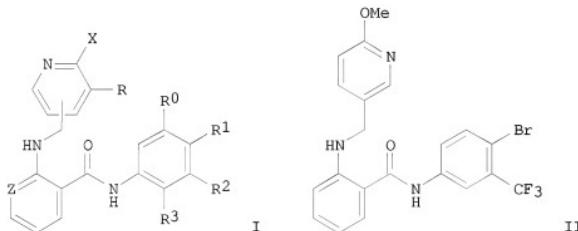
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DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:y

L7 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2010 ACS on STN
ACCESSION NUMBER: 2004:515506 HCAPLUS
DOCUMENT NUMBER: 141:71453
TITLE: Preparation of anthranilic acid amide derivatives as neoplastic inhibitors
INVENTOR(S): Bold, Guido; Furet, Pascal; Manley, Paul William
PATENT ASSIGNEE(S): Novartis Ag, Switz.; Novartis Pharma GmbH
SOURCE: PCT Int. Appl., 81 pp.
CODEN: PIXDZ
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
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| WO | 2004052884 | A1 | 20040624 | WO 2003-EP14086 | 20031211 |
| W: | AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
GE, GH, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
LT, LU, LV, MA, MD, MK, MN, MX, NI, NO, NZ, OM, PG, PH, PL, PT,
RO, RU, SC, SE, SG, SK, SY, TJ, TM, TN, TR, TT, UA, US, UZ, VC,
VN, YU, ZA, ZW | | | | |
| RW: | AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE,
DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE,
SI, SK, TR | | | | |
| CA | 2506164 | A1 | 20040624 | CA 2003-2506164 | 20031211 |
| AU | 2003294834 | A1 | 20040630 | AU 2003-294834 | 20031211 |
| EP | 1572686 | A1 | 20050914 | EP 2003-785795 | 20031211 |
| EP | 1572686 | B1 | 20090415 | | |
| R: | AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HK, SK | | | | |
| BR | 2003017292 | A | 20051108 | BR 2003-17292 | 20031211 |
| CN | 1720244 | A | 20060111 | CN 2003-80104845 | 20031211 |
| CN | 100427483 | C | 20081022 | | |
| JP | 2006511518 | T | 20060406 | JP 2004-558075 | 20031211 |
| AT | 428709 | T | 20090515 | AT 2003-785795 | 20031211 |
| PT | 1572686 | E | 20090714 | PT 2003-785795 | 20031211 |
| ES | 2324531 | T3 | 20090810 | ES 2003-785795 | 20031211 |
| US | 20060128684 | A1 | 20060615 | US 2005-538199 | 20050609 |
| RITY APPLN. INFO.: | | | | GB 2002-29022 | A 20021212 |
| | | | | WO 2003-EP14086 | W 20031211 |

OTHER SOURCE(S): MARPAT 141:71453
GI



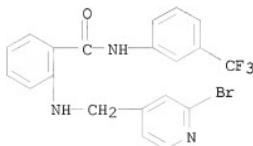
A8 The title compds. I (wherein R and R0 = independently H, halo, (un)substituted alkyl, alkenyl, alkynyl, aryl, heteroaryl, etc.; R1 = H, halo, (un)substituted alkyl, alkenyl, alkynyl, alkoxy, OCF3, OCH2CF3, OCH2CH2CF3, or OCH2CH2CH2CF3; R2 = perfluoroalkyl; R3 = H or halo; X = OH, alkoxy, alkylthio, imino, alkylimino, halo, etc.; Z = N or CH) or salts, N-oxides, or tautomers thereof are prepared as neoplastic inhibitors for the treatment of human or animal body. For example, the compound II was prepared in a multi-step synthesis. Formulations containing I as an active ingredient were also described.

IT 657401-06-4P

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RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(drug candidate, reactant; preparation of anthranilic acid amide derivs. as neoplastic inhibitors)

RN 657401-06-4 HCAPLUS
CN Benzanide, 2-[[((2-bromo-4-pyridinyl)methyl)amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2004120827 HCAPLUS

DOCUMENT NUMBER: 1401:181330

TITLE: Preparation of anthranylamidopyridines as inhibitors of vascular endothelial growth factor receptor-2 and -3 (VEGFR-2 and -3).

INVENTOR(S): Huth, Andreas; Krueger, Martin; Zorn, Ludwig; Ince, Stuart; Thierauch, Karl-Heinz; Menrad, Andreas; Haberey, Martin; Hess-Stump, Holger

PATENT ASSIGNEE(S): Schering Aktiengesellschaft, Germany
SOURCE: PCT Int. Appl., 88 pp.

DOCUMENT TYPE: Patent
LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|------------------|----------|
| WO 2004013102 | A1 | 20040212 | WO 2003-EP7964 | 20030722 |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
| RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG | | | | |
| DE 10235690 | A1 | 20040219 | DE 2002-10235690 | 20020731 |
| DE 10328036 | A1 | 20050105 | DE 2003-10328036 | 20030619 |
| CA 2493026 | A1 | 20040212 | CA 2003-2493026 | 20030722 |

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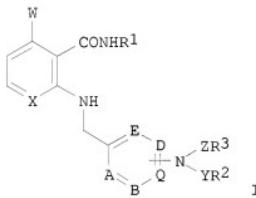
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| AU 2003281855 | A1 | 20040223 | AU 2003-281855 | 20030722 |
| BR 2003013122 | A | 20050705 | BR 2003-13122 | 20030722 |
| CN 1671666 | A | 20050921 | CN 2003-818334 | 20030722 |
| EP 1594841 | A1 | 20051116 | EP 2003-740470 | 20030722 |
| R: AT, BE, CH,
IE, SI, LT,
JP 2005538112 | DE, DK, ES, FR,
LV, FI, RO, MK, | GB, GR, IT, LI, LU,
CY, AL, TR, BG, CZ,
EE, HU, SK | MC, PT,
JP 2004-525272 | 20030722 |
| NZ 537291 | A | 20070223 | NZ 2003-537291 | 20030722 |
| US 20040147535 | A1 | 20040729 | US 2003-631018 | 20030731 |
| US 7148357 | B2 | 20061212 | | |
| US 20050054654 | A1 | 20050310 | US 2004-870491 | 20040618 |
| US 7517894 | B2 | 20090414 | | |
| MX 2004012948 | A | 20050912 | MX 2004-12948 | 20041217 |
| IN 2005DN00309 | A | 20070119 | IN 2005-DN309 | 20050127 |
| ZA 2005001642 | A | 20090930 | ZA 2005-1642 | 20050224 |
| NO 2005001035 | A | 20050429 | NO 2005-1035 | 20050225 |
| HR 2005000187 | A2 | 20051031 | HR 2005-187 | 20050225 |
| US 20070015794 | A1 | 20070118 | US 2006-525091 | 20060922 |
| US 7615565 | B2 | 20091110 | | |

PRIORITY APPLN. INFO.:

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| DE 2002-10235690 | A 20020731 |
| DE 2003-10328036 | A 20030619 |
| US 2002-407970P | P 20020905 |
| US 2003-483896P | P 20030702 |
| WO 2003-EP7964 | W 20030722 |
| US 2003-631018 | A3 20030731 |

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

OTHER SOURCE(S): MARPAT 140:181330
GI



AB Title compds. [I; X = CH, N; W = H, F; A, B, D, E, Q = N, C; ≤2 of A, B, D, E, Q = N; R1 = (substituted) aryl, heteroaryl; Y, Z = bond, CO, CS, SO2; R2, R3 = H, CONR9R10, SO2R6, COR11, NR9R10, (substituted) alkyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl; R2YNZR3 = atoms to form a 3-8 membered (substituted) (unsatd.) ring; R6 = H, alkyl, haloalkyl, (substituted) aryl, heteroaryl, NR9R10; R9, R10 = H, alkyl, alkenyl, aryl, cycloalkyl, etc.; R11 = alkyl, alkoxy, hydroxylalkyl, hydroxylalkoxy, cycloalkyl, (substituted) Ph, pyridyl, biphenyl, naphthyl], were prepared Thus, 2-[(2-bromopyridin-4-ylmethyl)amino]-N-(3-trifluoromethylphenyl)benzamide (preparation given) pyridine, and N,N-dimethylaminoethylamine were heated in a pressure vessel for 5 h at 200° to give 2-[(2-dimethylaminoethylamino)pyridin-4-

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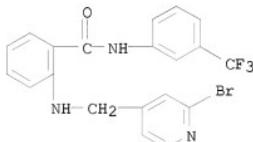
ylmethyl]amino]-N-(3-trifluoromethylphenyl)benzamide. It inhibited VEGFR-2 with IC₅₀ = 8-65 nM. It can be used for treatment of tumor or metastasis growth, psoriasis, Kaposi's sarcoma, restenosis, endometriosis, Crohn's disease, Hodgkin's disease, leukemia, arthritis, hemangioma, angiofibroma, eye disease, renal diseases, transplant rejection, fibrotic diseases, mesangial cell proliferative diseases, atherosclerosis, injuries to nervous tissue and for inhibition of the reocclusion of vessels after balloon catheter treatment, in vessel prosthetics, or after the application of mech. devices to hold open vessels, as immunosuppressants, for scar-free wound healing, age spots and contact dermatitis.

IT 657401-06-4

RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of anthranylamidopyridines as inhibitors of vascular endothelial growth factor receptor)

RN 657401-06-4 HCAPLUS

CN 2-[[{(2-bromo-4-pyridinyl)methyl]amino}-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



OS.CITING REF COUNT: 18 THERE ARE 18 CAPLUS RECORDS THAT CITE THIS RECORD (20 CITINGS)
REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2000:335388 HCAPLUS

DOCUMENT NUMBER: 132:347491

TITLE: Preparation of N-aryl(thio)anthranilic acid amides as VEGF receptor tyrosine kinase inhibitors

INVENTOR(S): Altmann, Karl-Heinz; Bold, Guido; Furet, Pascal; Manley, Paul William; Wood, Jeanette Marjorie; Ferrari, Stefano; Hofmann, Francesco; Mestan, Jurgen; Huth, Andreas; Kruger, Martin; Seidelmann, Dieter; Menrad, Andreas; Haberey, Martin; Thierauch, Karl-Heinz

PATENT ASSIGNEE(S): Novartis A.-G., Switz.; Novartis-Erfindungen Verwaltungsgesellschaft m.b.H.; Schering Aktiengesellschaft

SOURCE: PCT Int. Appl., 77 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.

KIND

APPLICATION NO.

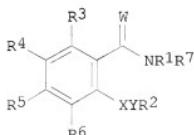
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| WO 2000027820 | A1 | 20000518 | WO 1999-EP8545 | 19991108 |
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| RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | | | | |
| CA 2346898 | A1 | 20000518 | CA 1999-2346898 | 19991108 |
| BR 9915210 | A | 20010724 | BR 1999-15210 | 19991108 |
| TR 200101237 | T2 | 20010821 | TR 2001-1237 | 19991108 |
| EP 1129075 | A1 | 20010905 | EP 1999-971802 | 19991108 |
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| HU 2001004188 | A2 | 20020328 | HU 2001-4188 | 19991108 |
| HU 2001004188 | A3 | 20020429 | | |
| JP 2002529453 | T | 20020910 | JP 2000-581000 | 19991108 |
| AU 758230 | B2 | 20030320 | AU 2000-13811 | 19991108 |
| NZ 511339 | A | 20030725 | NZ 1999-511339 | 19991108 |
| CN 1152014 | C | 20040602 | CN 1999-813108 | 19991108 |
| RU 2286338 | C2 | 20061027 | RU 2001-114978 | 19991108 |
| CZ 299829 | B6 | 20081210 | CZ 2001-1615 | 19991108 |
| SK 287259 | B6 | 20100407 | SK 2001-628 | 19991108 |
| NO 2001001894 | A | 20010704 | NO 2001-1894 | 20010417 |
| NO 328130 | B1 | 20091214 | | |
| ZA 2001003290 | A | 20030123 | ZA 2001-3290 | 20010423 |
| MX 2001004256 | A | 20030606 | MX 2001-4256 | 20010427 |
| US 20020019414 | A1 | 20020214 | US 2001-850434 | 20010507 |
| US 6448277 | B2 | 20020910 | | |
| IN 2001CN00638 | A | 20050304 | IN 2001-CN638 | 20010508 |
| ZA 2001004673 | A | 20020909 | ZA 2001-4673 | 20010607 |
| US 20030064992 | A1 | 20030403 | US 2002-180289 | 20020626 |
| US 6878720 | B2 | 20050412 | | |
| US 20040198782 | A1 | 20041007 | US 2004-828951 | 20040421 |
| US 7002022 | B2 | 20060221 | | |
| US 2006074112 | A1 | 20060406 | US 2005-254897 | 20051020 |
| PRIORITY APPLN. INFO.: | | | GB 1998-24579 | A 19981110 |
| | | | WO 1999-EP8545 | W 19991108 |
| | | | US 2001-850434 | A3 20010507 |
| | | | US 2002-180289 | A3 20020626 |
| | | | US 2004-828951 | A3 20040421 |

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

OTHER SOURCE(S): MARPAT 132:347491

GI



I

AB Use of title compds. I; W = O, S; X = NR8; Y = CR9R10(CH2)n, SO2; R9, R10 = H, alkyl; n = 0-3; R1 = aryl; R2 = mono- or bicyclic heteroaryl with the exception that R2 cannot = 2-phthalimidyl, and when Y = SO2 cannot represent 2,1,3-benzothiadiazol-4-yl; R3-R6 = H, substituent; R7, R8 = H, alkyl; or a N-oxide or a pharmaceutically acceptable salt thereof, for the preparation of a pharmaceutical product for the treatment of a neoplastic disease which responds to an inhibition of the VEGF receptor tyrosine kinase activity is claimed. Thus, a mixture of 4-pyridinecarboxaldehyde and 2-amino-N-(4-trifluoromethylphenyl)benzamide (preparation given) in MeOH containing

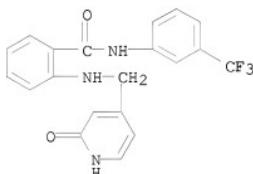
HOAc was treated with NaBH3CN followed by 16 h stirring to give 2-[(4-pyridyl)methyl]amino-N-[4-(trifluoromethyl)phenyl]benzamide. Tested I inhibited Flt-1 VEGF receptor tyrosine kinase with IC50 = 0.18-0.56 μM.

IT 269391-01-7P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of N-aryl(thio)anthranilic acid amides as VEGF receptor tyrosine kinase inhibitors)

RN 269391-01-7 HCPLUS

CN Benzamide, 2-[(1,2-dihydro-2-oxo-4-pyridinyl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



| | | |
|----------------------|----|---|
| OS.CITING REF COUNT: | 38 | THERE ARE 38 CAPLUS RECORDS THAT CITE THIS RECORD (42 CITINGS) |
| REFERENCE COUNT: | 14 | THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT |

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|--|--|------------|---------|
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| FULL ESTIMATED COST | | ENTRY | SESSION |
| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | | SINCE FILE | TOTAL |
| CA SUBSCRIBER PRICE | | ENTRY | SESSION |
| | | -2.55 | -2.55 |

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 3 JUN 2010 HIGHEST RN 1226953-63-4
DICTIONARY FILE UPDATES: 3 JUN 2010 HIGHEST RN 1226953-63-4

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stnqen/stndoc/properties.html>

=>
Uploading C:\Documents and Settings\brobinson1\My Documents\e-Red Folder\es6s6s6.str

1.10 STRUCTURE UPLOADED

=> s 110
SAMPLE SEARCH INITIATED 16:31:29 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1700 TO ITERATE

100.0% PROCESSED 1700 ITERATIONS 0 ANSWERS
SEARCH TIME: 00:00:01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
PROJECTED ITERATIONS: 31527 TO 36473
PROJECTED ANSWERS: 0 TO 0

1-11 0 SEA SSS SAM 1-10

```
=> s 110 full
THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 191.05 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y
FULL SEARCH INITIATED 16:31:33 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 33645 TO ITERATE
```

100.0% PROCESSED 33645 ITERATIONS 2 ANSWERS
SEARCH TIME: 00:00:01

L12 2 SEA SSSS FII: L10

$\Rightarrow d$ his

(FILE 'HOME' ENTERED AT 16:22:44 ON 05 JUN 2010)

stnvrksp

```
FILE 'REGISTRY' ENTERED AT 16:22:51 ON 05 JUN 2010
L1           STRUCTURE uploaded
L2           0 S L1
L3           0 S L1 FULL
L4           STRUCTURE uploaded
L5           0 S L4
L6           2 S L4 FULL
```

FILE 'HCAPLUS' ENTERED AT 16:27:33 ON 05 JUN 2010
L7 3 S L6
L8 0 S L7 AND LIGHTNER, J?/AU
L9 0 S L7 AND NG, H?/AU

FILE 'REGISTRY' ENTERED AT 16:29:11 ON 05 JUN 2010
L10 STRUCTURE uploaded
L11 0 S L10
L12 2 S L10 FULL

```

=> file hcaplus
COST IN U.S. DOLLARS                               SINCE FILE      TOTAL
                                                    ENTRY        SESSION
FULL ESTIMATED COST                           193.01       605.41

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)      SINCE FILE      TOTAL
                                                    ENTRY        SESSION
CA SUBSCRIBER PRICE                            0.00        -2.55

```

FILE 'HCAPLUS' ENTERED AT 16:31:43 ON 05 JUN 2010
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FILE COVERS 1907 - 5 Jun 2010 VOL 152 ISS 24
FILE LAST UPDATED: 4 Jun 2010 (20100604/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Apr 2010
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Apr 2010

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2010.

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This file contains CAS Registry Numbers for easy and accurate identification.

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substance identification.

=> s l12
L13 3 L12

=> d his

(FILE 'HOME' ENTERED AT 16:22:44 ON 05 JUN 2010)

FILE 'REGISTRY' ENTERED AT 16:22:51 ON 05 JUN 2010
L1 STRUCTURE uploaded
L2 0 S L1
L3 0 S L1 FULL
L4 STRUCTURE uploaded
L5 0 S L4
L6 2 S L4 FULL

FILE 'HCAPLUS' ENTERED AT 16:27:33 ON 05 JUN 2010
L7 3 S L6
L8 0 S L7 AND LIGHTNER, J?/AU
L9 0 S L7 AND NG, H?/AU

FILE 'REGISTRY' ENTERED AT 16:29:11 ON 05 JUN 2010
L10 STRUCTURE uploaded
L11 0 S L10
L12 2 S L10 FULL

FILE 'HCAPLUS' ENTERED AT 16:31:43 ON 05 JUN 2010
L13 3 S L12

=> s l13 not l7
L14 0 L13 NOT L7